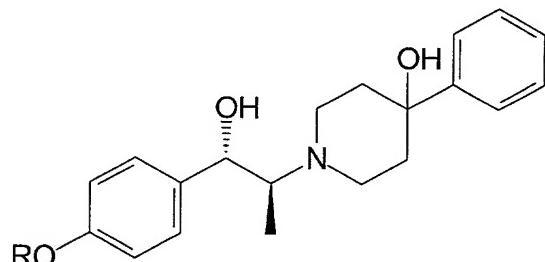


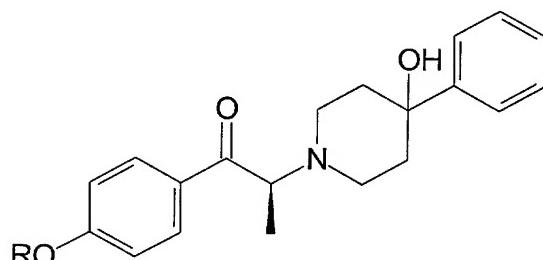
**CLAIMS:**

1        1. A process for the preparation of a nonracemic diastereomer selected  
2 from 1-(4-hydroxy-phenyl)-2-(4-hydroxy-4-phenyl-piperidin-1-yl)-1-propanol compounds of  
3 the structural formula I and stereoisomers thereof,



I

4  
5 wherein R is selected from hydrogen and hydroxyl protecting groups, comprising  
6 hydrogenating a corresponding nonracemic ketone selected from 1-(4-hydroxy-phenyl)-2-(4-  
7 hydroxy-4-phenyl-piperidin-1-yl)-1-propanone compounds of the structural formula II and  
8 enantiomers thereof,



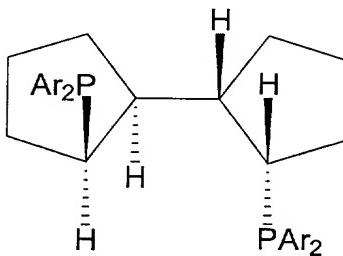
II

100-222-C-29

10        10 in the presence of a catalyst system comprising ruthenium, a nonracemic diphosphine ligand,  
11 a bidentate amine ligand selected from amino-thioethers and achiral diamines, and a base.

1        2. The process of claim 1 wherein the nonracemic diphosphine ligand  
2 comprises a 2,2'-bis(diorganophosphino)-1,1'-bis(cyclic) structure.

1        3. The process of claim 2 wherein the nonracemic diphosphine ligand is  
2 selected from enantiomers of diphosphine ligands having the structural formula



3  
4 wherein Ar is an aryl group.

- 1                  4.     The process of claim 3 wherein Ar is phenyl.
- 1                  5.     The process of claim 1 wherein the bidentate amine ligand is an amino-  
2     thioether.
- 1                  6.     The process of claim 5 wherein the amino-thioether is a  
2     2-(alkylthio)aniline.
- 1                  7.     The process of claim 6 wherein the 2-(alkylthio)aniline is selected  
2     from 2-(methylthio)aniline and 2-(ethylthio)aniline.
- 1                  8.     The process of claim 1 wherein the bidentate amine ligand is an achiral  
2     diamine.
- 1                  9.     The process of claim 8 wherein the achiral diamine comprises no chiral  
2     carbon centers.
- 1                  10.    The process of claim 8 wherein the achiral diamine is a 1,2-phenylene-  
2     diamine.
- 1                  11.    The process of claim 1 wherein the base is selected from basic  
2     inorganic and organic salts, alkylguanidines, aminophosphazenes, and proazaphosphatrane.
- 1                  12.    The process of claim 11 wherein the base is selected from  
2     alkylguanidines, aminophosphazenes, and proazaphosphatrane.
- 1                  13.    The process of claim 12 wherein the base is an alkylguanidine.
- 1                  14.    The process of claim 13 wherein the base is a pentaalkylguanidine.
- 1                  15.    The process of claim 1 wherein the hydroxyl protecting group is  
2     benzyl.
- 1                  16.    The process of claim 15 wherein the diastereomer is a *syn*-  
2     diastereomer.
- 1                  17.    The process of claim 16 wherein the *syn*-diastereomer is the (1*S*,2*S*)  
2     diastereomer.

1           **18.**   The process of claim **16** wherein the *syn*-diastereomer is formed in at  
2 least about 90% diastereomeric excess.

1           **19.**   A process for the preparation of (1*S*,2*S*)-1-(4-benzoxy-phenyl)-2-(4-  
2 hydroxy-4-phenyl-piperidin-1-yl)-1- by catalytic hydrogenation of (2*S*)-1-(4-benzyl-phenyl)-  
3 2-(4-hydroxy-4-phenyl-piperidin-1-yl)-1-propanone using a catalyst system comprising  
4 ruthenium, a (*S,S,S,S*)-2,2'-bis-(diarylphosphino)-1,1'-dicyclopentane ligand, a 1,2-phenylene  
5 diamine ligand, and a base.

1           **20.**   A process for the preparation of (1*S*,2*S*)-1-(4-benzoxy-phenyl)-2-(4-  
2 hydroxy-4-phenyl-piperidin-1-yl)-1- by catalytic hydrogenation of (2*S*)-1-(4-benzyl-phenyl)-  
3 2-(4-hydroxy-4-phenyl-piperidin-1-yl)-1-propanone using a catalyst system comprising  
4 ruthenium, a (*S,S,S,S*)-2,2'-bis-(diarylphosphino)-1,1'-dicyclopentane ligand, a  
5 2-(alkylthio)aniline ligand, and a base.